

CLAIMS

What is claimed is:

- 5 1. An implantable endocardial lead having a longitudinal axis and
extending between proximal and distal ends for use with a cardiac
stimulation device, the lead comprising:
- 10 an electrical conductor within the lead extending between
proximal and distal ends;
- an active fixation electrode comprising an electrically active
helix coaxial with the endocardial lead, coupled to the distal end of
the electrical conductor, and movable between a retracted position
fully within the lead and an extended position advanced beyond the
distal end of the lead for effecting penetration into the myocardial
15 tissue; and
- a guide system located proximally of the active fixation
electrode for rotating the electrically active helix about the
longitudinal axis as the helix is moved between the retracted and
extended positions.
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2. An implantable endocardial lead having a longitudinal axis and
extending between proximal and distal ends for use with a cardiac
stimulation device, the lead comprising:
- 25 an electrically active housing comprising a tubular end
region extending to a terminal rim at the distal end of the lead;
- an electrical conductor within the lead extending between
proximal and distal ends;
- an active fixation electrode within and spaced from the
electrically active housing and comprising an electrically active
30 helix coaxial with the endocardial lead coupled to the distal end of
the electrical conductor and movable between a retracted position
fully within the housing and an extended position advanced beyond

the terminal rim of the housing for effecting penetration into the myocardial tissue; and

a guide system located proximally of the active fixation electrode for rotating the electrically active helix about the longitudinal axis as the helix is moved between the retracted and extended positions.

3. An implantable endocardial lead as set forth in claim 2

wherein the electrically active housing comprises:

a generally planar bulkhead member extending transversely of the longitudinal axis; and

a tubular end region extending away from the bulkhead member to the terminal rim at the distal end of the lead; and

wherein the guide system comprises:

a spiral track member extending proximally away from the bulkhead member to a proximal rim distant from the bulkhead member; and

wherein the active fixation electrode comprises:

a conductive shaft having an outer peripheral surface and extending between proximal and distal ends and having an outwardly projecting follower member slidably engaged with the spiral track member, the electrical conductor being fixed to the proximal end thereof.

4. An implantable endocardial lead as set forth in claim 2 comprising:

an insulation sheath covering the electrical conductor, the sheath and the electrical conductor together defining an internal chamber extending from the proximal end to the distal end; and

an electrical connector being coupled to the proximal end of the electrical conductor.

5. An implantable endocardial lead as set forth in claim 2 and further comprising:

5 a resilient coupling mechanism for maintaining electrical continuity between the active fixation electrode and the electrically active housing throughout movement of the active fixation electrode between the retracted position and the extended position.

6. An implantable endocardial lead as set forth in claim 2

wherein the conductive shaft comprises:

10 an outer peripheral surface and extending between proximal and distal ends and having an outwardly projecting follower member slidably engaged with the spiral track member, the electrical conductor being fixed to the proximal end thereof;

15 an annular collar integral with the conductive shaft intermediate the proximal and distal ends and projecting radially from the longitudinal axis to an outer rim beyond the outer surface of the conductive shaft; and

20 a head portion coaxial with and extending distally from the annular collar and being of reduced diameter than the annular collar to define a distal annular shoulder at its intersection with the annular collar; and

wherein the spiral track member has an internal peripheral surface facing and slidably engaged with a part of the conductive shaft;

25 a compression spring intermediate and engaged with the bulkhead member and with the distal annular shoulder;

30 the annular collar being distant from the bulkhead member when the active fixation electrode is in the retracted position and being proximate the bulkhead member when the active fixation electrode is in the extended position, the compression spring biasing the annular collar in a direction away from the bulkhead member.

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7. An implantable endocardial lead as set forth in claim 2
wherein the active fixation electrode comprises an
electrically active helix advanceable outward relative to the distal
end of the conductor for effecting penetration into myocardial
tissue;

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8. An implantable endocardial lead as set forth in claim 2 and
further comprising:

a therapeutic element integral with the active fixation
electrode formed of a biocompatible matrix material being of
sufficient rigidity to penetrate the myocardial tissue.

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9. An implantable endocardial lead as set forth in claim 2 and
further comprising:

a therapeutic element generally cylindrical in shape coaxial
with and fixed on the distal end of the conductive shaft and formed
of a biocompatible matrix material being of sufficient rigidity to
penetrate the myocardial tissue.

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10. An implantable endocardial lead as set forth in claim 2
wherein the conductive coupling is cup-shaped and
comprises a base lying in a plane transverse of the longitudinal
axis of the conductive shaft and a cylindrical sidewall upstanding
from the base and coaxial with the longitudinal axis of the
conductive shaft, the base having a bore for slidable reception on
the conductive shaft; and

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wherein the compression spring extends between the base
and the conductive stopper.

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11. An implantable endocardial lead as set forth in claim 10
wherein the sidewall and base of the conductive coupling
define a cavity for reception of the conductive shaft, of the
conductive stopper, and of the compression spring coaxially
received on the conductive shaft.

12. An implantable endocardial lead as set forth in claim 2
wherein the active fixation electrode comprises an
electrically active helix advanceable outward relative to the distal
end of the conductor for effecting penetration into myocardial
tissue; and
wherein the electrically active housing comprises an
electrically active collar coaxial with the helix at the distal end of the
lead.

13. An implantable endocardial lead as set forth in claim 3
wherein the electrically active helix is fixed to the distal end
of the conductive shaft.

14. An implantable endocardial lead as set forth in claim 3
wherein the electrically active housing comprises:
a cylindrical guide member integral with and extending
proximally away from the bulkhead member and having an inner
facing peripheral surface; and
wherein the guide system comprises:
a spiral track member formed into the inner facing peripheral
surface of the cylindrical guide member and defined by opposed
spaced parallel side walls.

15. An implantable endocardial lead as set forth in claim 3

wherein the electrically active housing comprises:

a cylindrical guide member integral with and extending
proximally away from the bulkhead member and having an inner
facing peripheral surface; and

wherein the guide system comprises:

a spiral track member formed into the inner facing peripheral
surface of the cylindrical guide member and defined by opposed
spaced parallel side walls and a bottom wall connecting the side
walls.

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